Complete listing of claims:

- 1. (currently amended) A method of <u>improving the pharmacokinetic profile of</u> <u>administering</u> a drug for which the major clearance mechanism in humans is CYP2D6 mediated oxidative biotransformation, or a pharmaceutically acceptable salt thereof, comprising administering an <u>effective amount of the drug</u> in combination with a CYP2D6 inhibitor, or a pharmaceutically acceptable salt thereof, to a human in need of the intended pharmaceutical activity of such drug, wherein said drug and said CYP2D6 inhibitor are not the same compound, wherein said drug is (1S, 2S)-1-(4-hydroxyphenyl)-2-(4-hydroxy-4-phenylpiperidin-1-yl)-1-propanol or a pharmaceutically acceptable salt thereof.
- 2. (cancelled)
- 3. (cancelled)
- 4. (Original) A method according to claim 1, wherein the CYP2D6 inhibitor is quinidine, ajmalacine or pharmaceutically acceptable salts thereof.
- 5. (Original) A method according to claim 1, wherein the CYP2D6 inhibitor is selected from the group consisting of sertraline, venlafaxine, dexmedetomidine, tripennelamine, premethazine, hydroxyzine, halofrintane, chloroquine, moclobemide, and pharmaceutically acceptable salts thereof.
- 6. (Original) A method according to claim 1, wherein the CYP2D6 inhibitor is St. John's wort, or an extract of constituent thereof.